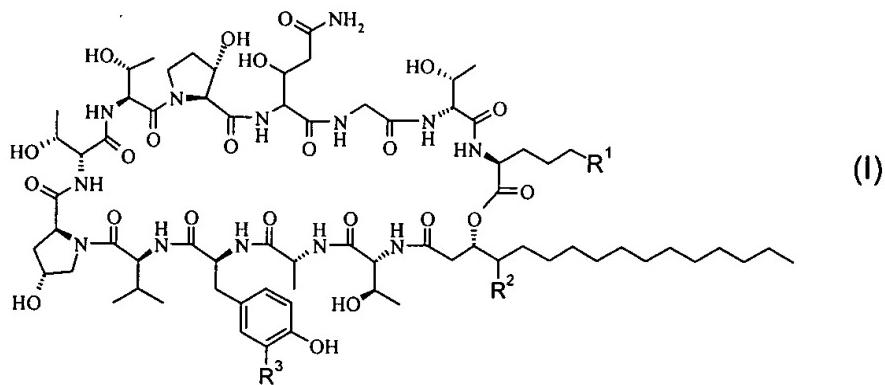


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Amendments to the Claims:

1. (Original) A compound of the formula,



wherein

R¹ is N-(3-aminopropyl)-N-[(2S)-2,5-diaminovaleryl]amino, N-(3-aminopropyl)-N-[-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, N-(3-aminopropyl)-N-[-5-amino-2-[N-(3-aminopropyl)amino]valeryl]amino, N-(2-aminoethyl)-N-[-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino or ornityl-ornitylamino;

R² is hydrogen or methyl;

R³ is hydrogen or hydroxy;

or a pharmaceutically acceptable salt thereof.

2. (Original) The compound according to Claim 1, wherein R¹ is N-(3-aminopropyl)-N-[(2S)-2,5-diaminovaleryl]amino, N-(3-aminopropyl)-N-[(2S)-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, N-(3-aminopropyl)-N-[(2R)-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, N-(3-aminopropyl)-N-[(2S)-5-amino-2-[N-(3-

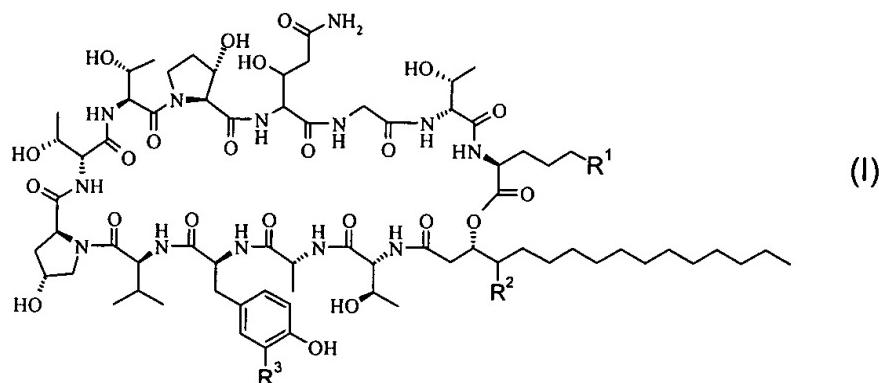
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aminopropyl)amino]valeryl]amino, N-(2-aminoethyl)-N-[(2S)-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, or (L)-ornityl-(D)-ornitylamino.

3. (Original) The compound according to Claim 1, wherein R² and R³ are hydrogen.
4. (Original) The compound according to Claim 1, wherein R¹ is N-(3-aminopropyl)-N-[(2S)-2,5-diaminovaleryl]amino, and R² and R³ are hydrogen.
5. (Original) The compound according to Claim 1, wherein R¹ is (L)-ornityl-(D)-ornitylamino, and R² and R³ are hydrogen.
6. (Original) The compound according to Claim 1, wherein R¹ is N-(3-aminopropyl)-N-[(2S)-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, and R² and R³ are hydrogen.
7. (Original) The compound according to Claim 1, wherein R¹ is N-(3-aminopropyl)-N-[(2R)-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, and R² and R³ are hydrogen.
8. (Original) The compound according to Claim 1, wherein R¹ is N-(3-aminopropyl)-N-[(2S)-5-amino-2-[N-(3-aminopropyl)amino]valeryl]amino, and R² and R³ are hydrogen.
9. (Original) The compound according to any of Claims 1 to 3, wherein R¹ is N-(2-aminoethyl)-N-[(2S)-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, and R² and R³ are hydrogen.
10. (Original) The compound according to Claim 1 wherein R² is methyl and R³ is hydrogen.
11. (Original) The compound according to Claim 1 wherein R² is hydrogen and R³ is hydroxyl.

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12. (Original) A pharmaceutical composition comprising a mixture of a pharmaceutically acceptable carrier and an effective amount of a compound of the formula



wherein

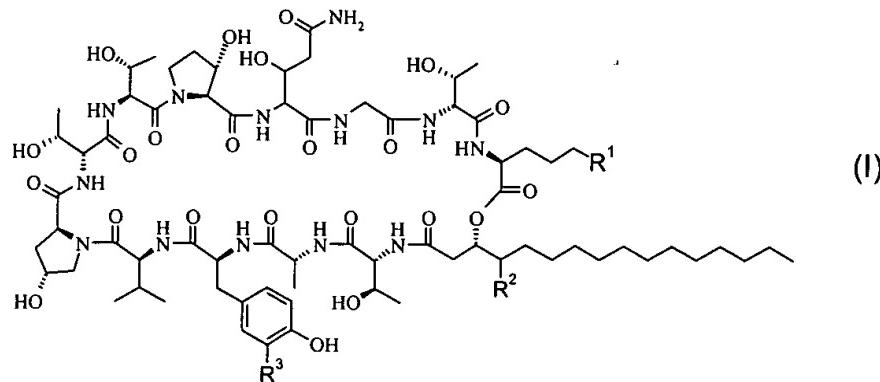
R^1 is N-(3-aminopropyl)-N-[(2S)-2,5-diaminovaleryl]amino, N-(3-aminopropyl)-N-[5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, N-(3-aminopropyl)-N-[5-amino-2-[N-(3-aminopropyl)amino]valeryl]amino, N-(2-aminoethyl)-N-[5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino or ornityl-ornitylamino;

R^2 is hydrogen or methyl;

R^3 is hydrogen or hydroxy;

or a pharmaceutically acceptable salt thereof.

13. (Currently amended) A method for the prophylactic or therapeutic treatment of mycoses which comprises administering to a human being or an animal an effective amount of the compound of the formula



wherein

R^1 is N-(3-aminopropyl)-N-[(2S)-2,5-diaminovaleryl]amino, N-(3-aminopropyl)-N-[-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino, N-(3-aminopropyl)-N-[-5-amino-2-[N-(3-aminopropyl)amino]valeryl]amino, N-(2-aminoethyl)-N-[-5-amino-2-[N,N-bis(2-aminoethyl)amino]valeryl]amino or ornityl-ornitylamino;

R^2 is hydrogen or methyl;

R^3 is hydrogen or hydroxy;

or a pharmaceutically acceptable salt thereof, thereby producing antifungal activity.